- b) contacting the peptide with a sample to be tested for the presence of the antibody;
- c) labeling antibody which binds to the peptide; and
- d) detecting the labeled antibody.

24 (New). The method of claim 23 wherein the labeled antibody in step (d) is detected by:

- a) labeling said antibody with a biotin-labeled Protein A/G conjugate;
- b) contacting the biotin-labeled Protein A/G conjugate with horse radish peroxidase conjugated streptavidin;
- c) contacting the horse radish peroxidase conjugated streptavidin with 3,3',5,5' tetramethyl benzidine (TMB); and
- d) detecting fluorescence.

25 (New). The method of Claim 23 wherein the sample is contacted with a plurality of peptides comprising an amino acid sequence selected from SEQ ID NOs: 1-7 and 9-15.

26 (New). The method of claim 23 wherein the sample is human serum.

27 (New). The method of claim 23 wherein the adenovirus is adenovirus 5.

Entry of the foregoing Amendment is requested. As of entry of the Amendment, claims 1-3, 8, 16, 18 and 21-27 will be pending. Each amended and new claim has written support in the specification; accordingly, no new matter has been added to the Application.

Written support for amended claim 1 appears in the specification, for example, at page 6, lines 8-25; at page 7, lines 6-30; at page 3, lines 24-25 and at claims 1-5, as filed.

Written support for amended claims 16 and 18 and new claims 21 and 22 appears in the specification, for example, at page 3, lines 1-8; at page 3, lines 12-20; at page 3, lines 24-25; at page 4, line 33 to page 5, line 3; at page 6, lines 8-25; at page 7, lines 6-30 and at claim 16, as filed.

Written support for new claims 23-27 appears in the specification, for example, at Example 1.9 (page 15, lines 8-28); page 5, lines 6-16; page 3, lines 24-25; and at page 4, line 30 to page 5, line 3.

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The changes to claims 2, 3 and 8 are formal changes which do not add any new matter; written support for these claims appears, for example, at the respective claims as filed.

Response to Notice to Comply with Requirements for Patent Applications Containing Nucleotide Sequence and/or Amino Acid Sequence Disclosures

A diskette is enclosed which includes a Substitute Sequence Listing for the above-referenced application which corrects the errors indicated in the Notice. A paper copy of the file is also attached.

The sequences in the present Substitute Sequence Listing were submitted with the original specification; therefore, no new matter has been added to the application. Furthermore, the contents of the attached paper entitled "SEQUENCE LISTING" and of the accompanying, identically labeled diskette, specifically the text file therein labeled "jb0976 seqlist.ST25", are the same.

Early and favorable action is earnestly solicited

Respectfully submitted

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IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicant(s): Mytych, et al.

Serial No.: 09/643,458

Filed: August 22, 2000

For: METHODS AND REAGENTS FOR THE

DETECTION OF ANTIBODIES TO ADENOVIRUS

Examiner: TBA

Group Art Unit: 1648

Confirmation No.: 9578

MARKED-UP AMENDMENT UNDER 37 C.F.R. §1.121

OCT 0 7 2002

Honorable Commissioner of Patents & Trademarks Washington, DC 20231

OFFICE OF PETITIONS

Sir:

The present Marked-Up Amendment shows all changes made to the claims in the enclosed Preliminary Amendment.

- 1 (Amended). A method for detecting <u>an antibody</u> [antibodies] capable of binding to adenovirus comprising <u>the steps of</u>:
- a) [immobilization of] <u>immobilizing</u> a peptide <u>comprising an amino acid sequence</u> <u>selected from SEQ ID NOs: 1-7 and 9-15</u> [capable of being bound by an anti-adenovirus antibody directly] onto a flowcell of a sensorchip in a biosensor;
- b) [obtaining a serum sample from a patient to be tested and contacting said serum] contacting a sample to be tested for the presence of the antibody with the immobilized peptide[,]; and

- c) [measuring] <u>detecting</u> binding of [antibodies] <u>the antibody</u> to the immobilized peptide by <u>detecting surface plasmon resonance in the biosensor</u> [means of response units from said biosensor].
- 2 (Amended). The method of Claim 1[,] wherein the adenovirus is [said peptide is capable of being bound by antibodies specific to] adenovirus 5.
- 3 (Amended). The method of Claim [2] 1, wherein said [serum] sample is human serum.
- 8 (Amended). The method of Claim 1 [further comprising] wherein a plurality of peptides comprising amino acid sequences selected from SEQ ID NOs: 1-7 and 9-15 [capable of being bound by an anti-adenovirus antibody] are directly immobilized, each on its own separate flowcell.
- 16 (Amended). A method for detecting an antibody [antibodies] capable of binding to adenovirus [,] comprising the steps of contacting a sample to be tested for the presence of the antibody with a peptide comprising an amino acid sequence selected from [the group consisting] SEQ ID NOs: 1-7 and 9-15 and detecting binding between the peptide and the antibody [SEQ ID NO: 1, SEQ ID NO: 2, SEQ ID NO: 3, SEQ ID NO: 4, SEQ ID NO: 5, SEQ ID NO: 6, and SEQ ID NO: 7 or a peptides having substantial sequence identity thereto].
- 18 (Amended). The method of Claim [17,] 16 [further comprising] wherein the sample is contacted with a plurality of peptides comprising an amino acid sequence selected from SEQ ID NOs: 1-7 and 9-15 [capable of being bound by an anti-adenovirus antibody].